



Should we use central venous saturation to guide management in high-risk surgical patients?

Pearse, RM; Hinds, CJ

- “The final publication is available at <http://ccforum.biomedcentral.com/articles/10.1186/cc5122>”

For additional information about this publication click this link.

<http://qmro.qmul.ac.uk/xmlui/handle/123456789/11479>

Information about this research object was correct at the time of download; we occasionally make corrections to records, please therefore check the published record when citing. For more information contact scholarlycommunications@qmul.ac.uk

Commentary

Should we use central venous saturation to guide management in high-risk surgical patients?

Rupert M Pearse and Charles J Hinds

Barts and The London School of Medicine and Dentistry, Queen Mary's University of London, 5th floor, 38 Little Britain, St. Bartholomew's Hospital, London EC1A 7BE, UK

Corresponding author: Rupert Pearse, rupert.pearse@bartsandthelondon.nhs.uk

Published: 15 December 2006

This article is online at <http://ccforum.com/content/10/6/181>

© 2006 BioMed Central Ltd

Critical Care 2006, **10**:181 (doi:10.1186/cc5122)

See related research by The Collaborative Study Group on Perioperative ScvO₂ Monitoring, <http://ccforum.com/content/10/6/R158>

Abstract

Measurements of central venous oxygen saturation (ScvO₂) have been successfully used to guide haemodynamic therapy in critical care. The efficacy of this approach in the treatment of severe sepsis and septic shock has stimulated interest in the use of ScvO₂ to guide management in patients undergoing major surgery. The physiological basis of ScvO₂ measurement is complex. A number of outstanding issues will need to be resolved before incorporating ScvO₂ measurement into routine practice. First, it is not yet clear which value of ScvO₂ should be targeted. Second, there is some uncertainty as to which interventions are the most effective for achieving the desired value of ScvO₂ or how long this value should be maintained. The study by The Collaborative Study Group on Perioperative ScvO₂ Monitoring published in this edition of *Critical Care* may help provide answers to some of these questions. Our understanding of ScvO₂ measurement remains limited, however, and the routine use of peri-operative ScvO₂-guided goal-directed therapy cannot be recommended until a large randomised trial has confirmed the value of this approach.

The use of central venous saturation (ScvO₂) to guide haemodynamic management is an important and evolving aspect of clinical practice. An observational study [1] published in this issue of *Critical Care* has advanced our understanding of this form of monitoring by exploring the association between derangements in ScvO₂ and complication rates after major abdominal surgery. This study provides a detailed description of peri-operative trends in ScvO₂ and confirms the findings of previous work which suggests that reductions in ScvO₂ are associated with increased post-operative complication rates [2]. Although the study is relatively small, the robust multi-centre approach and consistency with previous work support the applicability of the findings.

The comparative simplicity of ScvO₂ measurement makes this an attractive technique. With the blood gas analysis technology available in most institutions, intermittent ScvO₂

monitoring can be performed in any patient with a central venous catheter. However, it is not yet clear whether ScvO₂ measurement through intermittent blood sampling is an adequate alternative to continuous monitoring with a fibre-optic catheter. Interest in ScvO₂ measurement is not new, and several reports have explored the physiology and clinical significance of this parameter over the past 50 years [3]. Of these, the work of Rivers and colleagues [4] has proved the most influential. These authors used a ScvO₂ value of 70% as a target for goal-directed haemodynamic therapy (GDT) in patients presenting to hospital with severe sepsis and septic shock. They demonstrated that it may be possible to achieve substantial mortality reductions without the need for complex or invasive cardiac output monitoring technology. The success of Rivers' work and several trials of peri-operative GDT indicates that the use of ScvO₂ as a haemodynamic goal may be equally valuable in surgical patients [5-8].

However, several questions must be considered before embarking on an interventional trial of ScvO₂-guided peri-operative GDT. First, what treatments should be used to achieve the target value for ScvO₂? Second, which target value is most appropriate? Finally, how long should the target value be maintained? The study by the Collaborative Study Group (CSG) is important because it sets out to address some of these key questions. The value of ScvO₂ in any given patient reflects not only oxygen delivery but also oxygen consumption. Reductions in ScvO₂ may therefore reflect a large number of acute changes in physiology including hypoxia, shivering, anaesthesia, haemorrhage and myocardial ischaemia [3]. The therapeutic approach to achieving the target value may need to include more than simply intravenous fluids and inotropic therapy. If a period of post-operative sedation and invasive ventilation is required to control oxygen consumption, would such an intervention be

valid? Although the normal value of ScvO₂ is often quoted as 70%, there are in fact few published data to confirm this, either in healthy volunteers or in surgical patients [3]. Previous observational work shows that considerable variations in ScvO₂ may occur depending on the nature and severity of the acute physiological disturbance. It would be naive simply to accept this 'normal' value as being optimal in every clinical situation.

The CSG researchers explored the relationship between ScvO₂ and post-operative complication rates. Their findings suggest that a higher target value of 75% would be more appropriate in patients undergoing major abdominal surgery. This finding is consistent with the analysis of ScvO₂ data from a recent interventional trial of post-operative GDT [2]. However, both these studies have shown that large decreases in ScvO₂ occur immediately after surgery. It is unclear whether such changes, which are more marked in those patients who develop complications, relate predominantly to an increase in oxygen consumption, a decrease in oxygen delivery or, more probably, a failure to increase delivery to match increased consumption. What is more, these observations raise the possibility that the most appropriate goal for ScvO₂ may vary during and after surgery. The question of how long GDT should be continued remains unanswered. Several recent successful GDT trials have opted for short periods of early treatment lasting between 4 and 8 hours [4,6,7]. However, GDT has also been effective when administered for periods of up to 24 hours [5,8].

As with any monitoring technology, ScvO₂ is a double-edged sword. Anecdotal evidence suggests that clinicians have a limited understanding of the pitfalls associated with ScvO₂ measurement, which may lead to a number of problems in practice. For example, the aggressive targeting of too high a value for ScvO₂ may be harmful, particularly in the elderly. The authors make an important point in suggesting that the targeted value for ScvO₂ should be modified for different patient groups. In particular, the presence of cytopathic hypoxia in septic patients may result in a high value of ScvO₂ despite low oxygen delivery. Another consideration is that of sampling site. Venous oxygen saturation differs between the superior vena cava and the right atrium, and the value of ScvO₂ may therefore vary according to the position of the catheter tip [3]. Despite the promising findings of this most recent work, the routine peri-operative use of ScvO₂-guided GDT cannot be recommended until a large randomised trial has confirmed the value of this approach.

Competing interests

The authors declare that they have no competing interests.

References

1. The Collaborative Study Group on Perioperative ScvO₂ Monitoring: **Multicentre study on peri- and postoperative central venous oxygen saturation in high-risk surgical patients.** *Crit Care* 2006, **10**:R158.
2. Pearse RM, Dawson D, Fawcett J, Rhodes A, Grounds RM, Bennett ED: **Changes in central venous saturation after major surgery, and association with outcome.** *Crit Care* 2005, **9**:R694-R699.
3. Pearse RM, Rhodes A: **Mixed and central venous oxygen saturation.** In *Yearbook of Intensive Care and Emergency Medicine*. Edited by Vincent JL. Berlin: Springer; 2005:592-602.
4. Rivers E, Nguyen B, Havstad S, Ressler J, Muzzin A, Knoblich B, Peterson E, Tomlanovich M: **Early goal-directed therapy in the treatment of severe sepsis and septic shock.** *N Engl J Med* 2001, **345**:1368-1377.
5. Boyd O, Grounds RM, Bennett ED: **A randomized clinical trial of the effect of deliberate perioperative increase of oxygen delivery on mortality in high-risk surgical patients.** *JAMA* 1993, **270**:2699-2707.
6. McKendry M, McGloin H, Saberi D, Caudwell L, Brady AR, Singer M: **Randomised controlled trial assessing the impact of a nurse delivered, flow monitored protocol for optimisation of circulatory status after cardiac surgery.** *BMJ* 2004, **329**:258.
7. Pearse RM, Dawson D, Fawcett J, Rhodes A, Grounds RM, Bennett ED: **Early goal-directed therapy after major surgery reduces complications and duration of hospital stay. A randomised, controlled trial.** *Crit Care* 2005, **9**:R687-R693.
8. Polonen P, Ruokonen E, Hippelainen M, Poyhonen M, Takala J: **A prospective, randomized study of goal-oriented hemodynamic therapy in cardiac surgical patients.** *Anesth Analg* 2000, **90**:1052-1059.
9. Gattinoni L, Brazzi L, Pelosi P, Latini R, Tognoni G, Pesenti A, Antonio, Fumagalli, Roberto, The SvO₂ Collaborative Group: **A trial of goal-oriented hemodynamic therapy in critically ill patients.** *N Engl J Med* 1995, **333**:1025-1032.
10. Hayes MA, Timmins AC, Yau EH, Palazzo M, Hinds CJ, Watson D: **Elevation of systemic oxygen delivery in the treatment of critically ill patients.** *N Engl J Med* 1994, **330**:1717-1722.